Application No. 08/203,004 Docket No. 061266-5001-03 Amendment dated July 9, 2008 Response to Office action mailed January 9, 2008

## Amendments to the Claims:

This listing of claims will replace all prior versions, and listings of claims in the application.

## Listing of Claims:

- 1.-46. (canceled)
- 47. (currently amended) A method of treating a malignant tumor in a human patient comprising co-administering to the patient
- (a) a composition comprising a therapeutically effective amount of human tumor cells that:
  - (i) are conjugated to a hapten;
- (ii) are of the same tumor type as the malignant tumor of said patient for treatment of whom the composition is intended;
  - (iii) are autologous to said patient; and
- (iv) have been rendered incapable of growing in the body of a human upon injection therein; and
  - (b) an adjuvant;

wherein said malignant tumor is from a cancer selected from the group consisting of melanoma cancer, lung cancer, colon cancer, breast cancer, kidney cancer, and prostate cancer; and wherein said administration elicits a delayed-type hypersensitivity response by said patient to tumor cells conjugated to said hapten; and

repeating said administration of said composition for a total of at least six administrations of said composition; and

administering a therapeutically effective amount of cyclophosphamide to the patient only prior to the first administration of said composition.

Application No. 08/203,004 Docket No. 061266-5001-03 Amendment dated July 9, 2008 Response to Office action mailed January 9, 2008

48-66. (canceled)

- 67. (previously presented) The method of claim 47, wherein said hapten is selected from the group consisting of dinitrophenyl, trinitrophenyl, and N-iodoacetyl-N'-(5-sulfonic 1 -naphtyl) ethylene diamine.
  - 68. (previously presented) The method of claim 47 wherein said hapten is dinitrophenyl.
  - 69. (canceled)
- 70. (currently amended) The method of claim 47, wherein further comprising administering a therapeutically effective amount of cyclophosphamide comprises administering a dose of about 300 mg/M<sup>2</sup> of cyclophosphamide prior to the first administration of said composition.
  - 71. (canceled)
- 72. (currently amended) The method of claim 47 70 further comprising sensitizing the patient with a therapeutically effective amount of 1 -fluoro-2,4-dinitrobenzene prior to administering the thereapeutically effective amount of cyclophosphamide.
  - 73. (canceled)
- 74. (previously presented) The method of claim 47 wherein said adjuvant is Bacillus Calmette-Guerin.
- 75. (previously presented) The method of claim 47 wherein said administration of said composition prolongs survival of said patient.
  - 76. (canceled)
- 77. (previously presented) The method of claim 47, wherein said administration of said composition elicits T lymphocytes that infiltrate the tumor of said human.